**EXPLANATORY STATEMENT**

***National Health Act 1953***

***National Health (Listed drugs on F1 or F2) Amendment Determination 2020 (No. 8)***

**PB 84 of 2020**

**Authority**

This instrument, made under subsection 85AB(1) of the *National Health Act 1953* (the Act), amends the *National Health (Listed drugs on F1 or F2) Determination 2010* (PB 93 of 2010) (the Principal Determination).

The Principal Determination provides for the allocation of drugs to the F1 and F2 formularies of the Pharmaceutical Benefits Scheme (PBS).

**Purpose**

This instrument makes amendments to the Principal Determination.

The Act provides that listed drugs may be assigned to formularies identified as F1 and F2.

F1 is intended for single brand drugs and F2 for drugs that have multiple brands, or are in a therapeutic group with other drugs with multiple brands. Drugs on F2 are subject to the provisions of the Act relating to price disclosure and guarantee of supply.

Section 84AC of the Act provides that a drug is on F1 or F2 if there is a determination in force under section 85AB that the drug is on F1 or F2.

Subsection 85AB(1) of the Act empowers the Minister (or delegate) to determine by legislative instrument that a listed drug is on F1 or F2. For a drug to be on F1, it must satisfy the criteria in subsection 85AB(4). This requires that there are no listed brands of pharmaceutical items that have the drug that are bioequivalent or biosimilar, and no listed brands of pharmaceutical items that have another drug in the same therapeutic group as the first drug that are bioequivalent or biosimilar. It also requires that the drug was not on F2 the day before the determination comes into effect. A drug may only be determined to be on F2 if it does not satisfy one or more of the criteria for F1 (subsection 85AB(3)).

When subsection 85AB(5) of the Act applies, which relates to listed drugs with a single brand combination item on the PBS, the listed drug is not placed on F1 or F2, but on the administrative combination drug list.

This Instrument amends the Principal Determination by adding to F1 one new drug, acalabrutinib.

**Variation and revocation**

Unless there is an express power to revoke or vary PB 93 of 2010 cited in this Instrument, subsection 33(3) of the *Acts Interpretation Act 1901* is relied upon to revoke or vary PB 93 of 2010.

**Consultation**

This Instrument affects pharmaceutical companies with new medicines listing on the PBS. Acalabrutinib meets the criteria for F1 set out in section 85AB(4) of the Act.

Before this drug was listed and allocated to the F1 formulary, there were detailed consultations about the drug with the responsible person, and a recommendation was received from the Pharmaceutical Benefits Advisory Committee (PBAC). Any PBAC recommendation is made following receipt of a submission made by the affected pharmaceutical company. Two-thirds of the PBAC membership is from the following interests or professions: consumers, health economists, practising community pharmacists, general practitioners, clinical pharmacologists and medical specialists.

No additional consultations with experts was undertaken regarding this determination because consultation with the affected responsible person and the PBAC drew on the knowledge of persons with relevant expertise.

**Commencement**

This Instrument commences on 1 September 2020.

This Instrument constitutes a legislative instrument for the purpose of the  
*Legislation Act 2003*.

**Statement of Compatibility with Human Rights**

*Prepared in accordance with Part 3 of the Human Rights (Parliamentary Scrutiny) Act 2011*

***National Health (Listed Drugs on F1 or F2) Amendment Determination 2020 (No. 8)   
(PB 84 of 2020)***

This Legislative Instrument is compatible with the human rights and freedoms recognised or declared in the international instruments listed in section 3 of the *Human Rights (Parliamentary Scrutiny) Act 2011*.

**Overview of the Legislative Instrument**

This Legislative Instrument is made pursuant to subsection 85AB(1) of the *National Health Act 1953* (the Act), which relates to listed drugs on F1 or F2. This Instrument amends the *National Health (Listed drugs on F1 or F2) Determination 2010* (PB 93 of 2010) (the Principal Determination) which provides for the allocation of drugs to the F1 and F2 formularies of the Pharmaceutical Benefits Scheme (PBS).

This Instrument amends the Principal Determination by adding to F1 one new drug, acalabrutinib.

**Human rights implications**

This Legislative Instrument engages Article 2 and 12 of the International Covenant on Economic, Social and Cultural Rights (ICESCR) by assisting with the progressive realisation by all appropriate means of the right of everyone to the enjoyment of the highest attainable standard of physical and mental health.

The PBS is a benefit scheme which assists with advancement of this human right by providing for subsidised access by patients to medicines. The recommendatory role of the Pharmaceutical Benefits Advisory Committee (PBAC) ensures that decisions about subsidised access to medicines on the PBS are evidence-based.

**Conclusion**

This Legislative Instrument is compatible with human rights. Human rights continue to be protected by retaining on the PBS clinically important medicines and placing them in formularies that ensure the most cost effective pricing for supply of each medicine to Australians.

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